

LAXATIVE- bisacodyl tablet, delayed release
Publix Super Markets Inc

Publix 44-676A-Laxative

Active ingredient (in each tablet)

Bisacodyl USP, 5 mg

Purpose

Stimulant laxative

Uses

- for relief of occasional constipation (irregularity)
- this product generally produces bowel movement in 6 to 12 hours

Warnings

Do not use

if you cannot swallow without chewing.

Ask a doctor before use if you have

- a sudden change in bowel habits that lasts more than 2 weeks
- stomach pain, nausea or vomiting

When using this product

- do not chew or crush tablet(s)
- do not use within 1 hour after taking an antacid or milk
- it may cause stomach discomfort, faintness, and cramps

Stop use and ask a doctor if

- you have rectal bleeding or fail to have a bowel movement after use of a laxative. These could be signs of a serious condition.
- you need to use a laxative for more than 1 week

If pregnant or breast-feeding,

ask a health professional before use.

Keep out of reach of children.

In case of overdose, get medical help or contact a Poison Control Center right away.

Directions

- **do not take more than directed**
- take with a glass of water

adults and children 12 years and over	take 1 to 3 tablets in a single daily dose
children 6 to under 12 years	take 1 tablet in a single daily dose
children under 6 years	ask a doctor

Other information

- **TAMPER EVIDENT: DO NOT USE IF OUTER PACKAGE IS OPENED OR BLISTER IS TORN OR BROKEN**
- avoid excessive humidity
- store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F)
- see end flap for expiration date and lot number

Inactive ingredients

carmine, colloidal silicon dioxide, corn starch, FD&C red #40 aluminum lake, hypromellose, lactose anhydrous, magnesium stearate, methacrylic acid, microcrystalline cellulose, polydextrose, polyethylene glycol, shellac wax, simethicone, sodium bicarbonate, sodium lauryl sulfate, stearic acid, talc, titanium dioxide, triacetin, triethyl citrate

Principal display panel

NDC 56062-976-01

P

laxative

BISACODYL USP, 5 mg

STIMULANT LAXATIVE

- Gentle, dependable constipation relief

30

COMFORT

COATED TABLETS

ACTUAL SIZE

TAMPER EVIDENT: DO NOT USE IF PACKAGE IS OPENED OR IF BLISTER UNIT IS TORN, BROKEN OR SHOWS ANY SIGNS OF TAMPERING

50844 REV0923A67601

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B9991R1

Drug Facts (continued)

Warnings
Do not use if you cannot swallow without chewing.
Ask a doctor before use if you have a sudden change in bowel habits that lasts more than 2 weeks
stomach pain, nausea or vomiting

When using this product
do not chew or crush tablets
do not use within 1 hour after taking an antacid or milk
it may cause stomach discomfort, flatness, and cramps

Stop use and ask a doctor if
you have rectal bleeding or fail to have a bowel movement after use of a laxative. These could be signs of a serious condition
you need to use a laxative for more than 1 week
If pregnant or breast-feeding, ask a health professional before use.
Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away.

Drug Facts (continued)

Inactive ingredients
#40 aluminum lake, hypromellose, lactose anhydrous, carmine, colloidal silicon dioxide, corn starch, FD&C red magnesium stearate, methacrylic acid, microcrystalline

Other information
TAMPER EVIDENT: DO NOT USE IF OUTER PACKAGE IS OPENED OR BLISTER IS TORN OR BROKEN
avoid excessive humidity
store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F)
see end flap for expiration date and lot number

Directions do not take more than directed
take with a glass of water
adults and children 12 years take 1 to 3 tablets in a single daily dose
children 6 to under 12 years take 1 tablet in a single daily dose
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Drug Facts
KEEP OUTER PACKAGE FOR COMPLETE PRODUCT INFORMATION

Active ingredient (in each tablet)
Bisacodyl USP, 5 mg Stimulant laxative

Purpose
Stimulant laxative

Uses
for relief of occasional constipation (irregularity)
this product generally produces bowel movement in 6 to 12 hours

Drug Facts (continued)

B-1518-676A-01-R
REV0923A67601

NDC 56062-976-01

laxative
BISACODYL USP, 5 mg
STIMULANT LAXATIVE

● Gentle, dependable constipation relief

30 COMFORT COATED TABLETS


ACTUAL SIZE

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Drug Facts (continued)
cellulose, polyethylene glycol, shellac wax, simethicone, sodium bicarbonate, sodium lauryl sulfate, stearic acid, talc, titanium dioxide, thacebn, triethyl citrate

TAMPER EVIDENT: DO NOT USE IF PACKAGE IS OPENED OR IF BLISTER UNIT IS TORN, BROKEN OR SHOWS ANY SIGNS OF TAMPERING

No Print / No Varnish
Lot no. & Exp. date



LAXATIVE

bisacodyl tablet, delayed release

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:56062-976
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BISACODYL (UNII: 10X0709Y6I) (DEACETYLBISACODYL - UNII:R09078E41Y)	BISACODYL	5 mg

Inactive Ingredients

Ingredient Name	Strength
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
STARCH, CORN (UNII: O8232NY3SJ)	
FD&C RED NO. 40 ALUMINUM LAKE (UNII: 6T47AS764T)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
ANHYDROUS LACTOSE (UNII: 3SY5LH9PMK)	
MAGNESIUM STEARATE (UNII: 70097M6130)	
METHACRYLIC ACID (UNII: 1CS02G8656)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYDEXTROSE (UNII: VH2XOU12IE)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
SHELLAC (UNII: 46N107B710)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
WATER (UNII: 059QF0KO0R)	
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
TRIACETIN (UNII: XHX3C3X673)	
TRIETHYL CITRATE (UNII: 8Z96QXD6UM)	

Product Characteristics

Color	pink	Score	no score
Shape	ROUND	Size	8mm
Flavor		Imprint Code	B
Contains			

Packaging

		Marketing Start	Marketing End
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#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:56062-976-01	2 in 1 CARTON	05/21/2024	
1		15 in 1 BLISTER PACK; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	05/21/2024	

Labeler - Publix Super Markets Inc (006922009)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		038154464	pack(56062-976)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867837	manufacture(56062-976) , pack(56062-976)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867894	manufacture(56062-976)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		967626305	pack(56062-976)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		117025878	manufacture(56062-976)

Revised: 5/2025

Publix Super Markets Inc